ADVIL PM- diphenhydramine hcl, ibuprofen capsule, liquid filled Haleon US Holdings LLC

Drug Facts

Active ingredients (in each capsule)

Diphenhydramine hydrochloride 25 mg Solubilized ibuprofen equal to 200 mg ibuprofen (NSAID)* (present as the free acid and potassium salt) *nonsteroidal anti-inflammatory drug

Purposes

Nighttime sleep-aid

Pain reliever

Uses

- for relief of occasional sleeplessness when associated with minor aches and pains
- helps you fall asleep and stay asleep

Warnings

Allergy alert:

Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

- hives
- facial swelling
- asthma (wheezing)
- shock
- skin reddening
- rash
- blisters

If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning:

This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you

• are age 60 or older

- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Heart attack and stroke warning:

NSAIDs, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or for longer than directed.

Do not use

- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- unless you have time for a full night's sleep
- in children under 12 years of age
- right before or after heart surgery
- with any other product containing diphenhydramine, even one used on skin
- if you have sleeplessness without pain

Ask a doctor before use if

- stomach bleeding warning applies to you
- you have problems or serious side effects from taking pain relievers or fever reducers
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke.
- you are taking a diuretic
- you have a breathing problem such as emphysema or chronic bronchitis
- you have glaucoma
- you have trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

- taking sedatives or tranquilizers, or any other sleep-aid
- under a doctor's care for any continuing medical illness
- taking any other antihistamines
- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- taking any other drug

When using this product

- drowsiness will occur
- avoid alcoholic drinks
- do not drive a motor vehicle or operate machinery
- take with food or milk if stomach upset occurs

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
 - feel faint
 - vomit blood
 - have bloody or black stools
 - have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
 - chest pain
 - trouble breathing
 - weakness in one part or side of body
 - slurred speech
 - leg swelling
- pain gets worse or lasts more than 10 days
- sleeplessness persists continuously for more than 2 weeks. Insomnia may be a symptom of a serious underlying medical illness.
- redness or swelling is present in the painful area
- any new symptoms appear

If pregnant or breast-feeding,

ask a health professional before use. It is especially important not to use ibuprofen at 20 weeks or later in pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- do not take more than directed
- adults and children 12 years and over: take 2 capsules at bedtime
- do not take more than 2 capsules in 24 hours

Other information

- each capsule contains: potassium 20 mg
- read all warnings and directions before use. Keep carton.
- store at 20-25°C (68-77°F)
- avoid excessive heat above 40°C (104°F)
- protect from light

Inactive ingredients

D&C red no. 33, FD&C blue no. 1, gelatin, medium-chain triglycerides, pharmaceutical ink, polyethylene glycol, potassium hydroxide, purified water, sorbitol sorbitan solution

Questions or Comments?

Call weekdays 9 AM to 5 PM EST at 1-800-88-ADVIL

Additional Information

Do Not Use if seal under bottle cap imprinted with "SEALED for YOUR PROTECTION" is broken or missing.

For most recent product information, visit www.Advil.com

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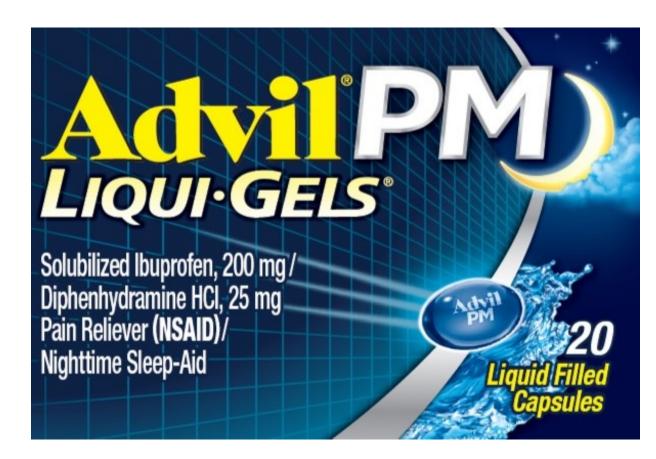
of Catalent Pharma Solutions.

PRINCIPAL DISPLAY PANEL (Advil PM Liqui-Gels)

Advil[®] PM LIQUI•GELS[®]

Solubilized Ibuprofen, 200 mg / Diphenhydramine HCl, 25 mg Pain Reliever **(NSAID)**/ Nighttime Sleep-Aid

20 Liquid Filled Capsules



Principal Display Panel (Advil PM Liqui-Gels Minis)

Advil® PM LIQUI•GELS®minis

NEWSmaller Capsule

Same Strength*

Solubilized Ibuprofen, 200 mg / Diphenhydramine HCl, 25 mg Pain Reliever **(NSAID)**/ Nighttime Sleep-Aid

*Compared to Advil PM Liqui-Gels

40 Liquid Filled Capsules



ADVIL PM

diphenhydramine hcl, ibuprofen capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0573-0167
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII: WK2XYI10QM)	IBUPROFEN	200 mg

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T)	
WATER (UNII: 059QF0KO0R)	
SORBITAN (UNII: 6092ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

Product Charact	eristics		
Color	BLUE (Clear blue)	Score	no score
Shape	OVAL (Oval liqui-gel)	Size	15mm
Flavor		Imprint Code	Advil;PM
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0573- 0167-10	16 in 1 CARTON	12/20/2005	12/07/2017
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0573- 0167-20	32 in 1 CARTON	12/20/2005	12/07/2017
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:0573- 0167-21	40 in 1 CARTON	12/20/2005	12/07/2017
3		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0573- 0167-25	1 in 1 CARTON	12/20/2005	
4		20 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:0573- 0167-43	1 in 1 CARTON	12/20/2005	
5		40 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:0573- 0167-55	1 in 1 CARTON	12/20/2005	
6		80 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:0573- 0167-18	24 in 1 CARTON	12/20/2005	12/07/2017
7		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:0573- 0167-19	24 in 1 CARTON	12/20/2005	12/07/2017
8		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
9	NDC:0573- 0167-26	1 in 1 CARTON	12/20/2005	
9		30 in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:0573- 0167-42	40 in 1 CARTON	12/20/2005	12/07/2017
10		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
11	NDC:0573- 0167-44	1 in 1 CARTON	12/20/2005	
11		50 in 1 BOTTLE; Type 0: Not a Combination Product		
12	NDC:0573- 0167-56	1 in 1 CARTON	12/20/2005	

12	100 in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing	Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021393	12/20/2005	

ADVIL PM

diphenhydramine hcl, ibuprofen capsule, liquid filled

Product Info	mation					
Product Type		HUMAN OTC DRUG	Item Code (S	Source)	NDC:057	3-0148
Route of Admin	istration	ORAL				
Active Ingred	ient/Active	Moiety				
	Ingree	dient Name		Basis of S	trength	Strength
DIPHENHYDRAMINE (DIPHENHYDRAMINE		DRIDE (UNII: TC2D6JAD40 83M))	DIPHENHYDRAM HYDROCHLORID		25 mg
IBUPROFEN (UNII:	WK2XYI10QM) (I	BUPROFEN - UNII:WK2XYI1	.0QM)	IBUPROFEN		200 mg
Inactive Ingre	edients					
		Ingredient Name			S	trength
D&C RED NO. 33	(UNII: 9DBA0SB	30L)				
FD&C BLUE NO. 1	L (UNII: H3R47K3	TBD)				
GELATIN, UNSPEC	CIFIED (UNII: 2G	86QN327L)				
MEDIUM-CHAIN T	RIGLYCERIDES	(UNII: C9H2L21V7U)				
POLYETHYLENE G	GLYCOL, UNSPE	CIFIED (UNII: 3WJQ0SDW	/1А)			
POTASSIUM HYDE	ROXIDE (UNII: W	ZH3C48M4T)				
WATER (UNII: 0590	QF0KO0R)					
SORBITAN (UNII: 6	O92ICV9RU)					
SORBITOL (UNII: 5	06T60A25R)					
Product Char	acteristics					
Color	BLUE (Clear	blue)	Score	n	o score	
Shape	OVAL (Oval li	qui-gel)	Size		3mm	
Flavor			Imprint Code	A	dvil;PM;Minis	;
Contains						
Packaging						
			Morte	eting Start	Marko	ting End

м	larketing l	nformation		
Μ	larketing l	nformation		
Μ	Marketing	Application Number or Monograph	Marketing Start	-
Μ	•		Marketing Start Date	Marketing En Date

Labeler - Haleon US Holdings LLC (079944263)

Revised: 1/2025

Haleon US Holdings LLC